

lead to a rule (1) having an annual effect on the economy of \$100 million or more, or adversely and materially affecting a sector of the economy, productivity, competition, jobs, the environment, public health or safety, or State, local or tribal governments or communities (also known as "economically significant"); (2) creating serious inconsistency or otherwise interfering with an action taken or planned by another agency; (3) materially altering the budgetary impacts of entitlement, grants, user fees, or loan programs; or (4) raising novel legal or policy issues arising out of legal mandates, the President's priorities, or the principles set forth in this Executive Order.

Pursuant to the terms of this Executive Order, EPA has determined that this rule is not "significant" and is therefore not subject to OMB review.

Pursuant to the requirements of the Regulatory Flexibility Act (Pub. L. 96-354, 94 Stat. 1164, 5 U.S.C. 601-612), the Administrator has determined that regulations establishing new tolerances or raising tolerance levels or establishing exemptions from tolerance requirements do not have a significant economic impact on a substantial number of small entities. A certification statement to this effect was published in the **Federal Register** of May 4, 1981 (46 FR 24950).

#### List of Subjects in 40 CFR Part 180

Environmental protection, Administrative practice and procedure, Agricultural commodities, Pesticides and pests, Reporting and recordkeeping requirements.

Dated: July 31, 1995.

**Daniel M. Barolo,**

*Director, Office of Pesticide Programs.*

Therefore, chapter I of title 40 of the Code of Federal Regulations is amended as follows:

#### PART 180—[AMENDED]

##### 1. In part 180:

a. The authority citation for part 180 continues to read as follows:

**Authority:** 21 U.S.C. 346a and 371.

b. By adding new § 180.484, to read as follows:

#### § 180.484 Flutolanil (N-(3-(1-methylethoxy)phenyl)-2-(trifluoromethyl)benzamide); tolerances for residues.

Tolerances are established for residues of flutolanil, N-(3-(1-methylethoxy)phenyl)-2-(trifluoromethyl)benzamide, and its metabolites converted to 2-

(trifluoromethyl) benzoic acid and calculated as flutolanil in or on the following raw agricultural commodities:

Commodity	Parts per million
Cattle, fat .....	0.10
Cattle, kidney .....	1.00
Cattle, liver .....	2.00
Cattle, mbyp .....	0.05
Cattle, meat .....	0.05
Cattle, milk .....	0.05
Eggs .....	0.05
Goats, fat .....	0.10
Goats, kidney .....	1.00
Goats, liver .....	2.00
Goats, mbyp .....	0.05
Goats, meat .....	0.05
Goats, milk .....	0.05
Hogs, fat .....	0.10
Hogs, kidney .....	1.00
Hogs, liver .....	2.00
Hogs, mbyp .....	0.05
Hogs, meat .....	0.05
Hogs, milk .....	0.05
Horses, fat .....	0.10
Horses, kidney .....	1.00
Horses, liver .....	2.00
Horses, mbyp .....	0.05
Horses, meat .....	0.05
Horses, milk .....	0.05
Peanuts .....	0.5
Peanut hay .....	15.0
Peanut hulls .....	5.0
Poultry (including turkeys), fat ..	0.05
Poultry (including turkeys), mbyp .....	0.05
Poultry (including turkeys), meat .....	0.05
Sheep, fat .....	0.10
Sheep, kidney .....	1.00
Sheep, liver .....	2.00
Sheep, meat .....	0.05
Sheep, mbyp .....	0.05
Sheep, milk .....	0.05

#### PART 185—[AMENDED]

##### 2. In part 185:

a. The authority citation for part 185 continues to read as follows:

**Authority:** 21 U.S.C. 346a and 348.

b. By adding new § 185.3385, to read as follows:

#### § 185.3385 Flutolanil (N-(3-(1-methylethoxy)phenyl)-2-(trifluoromethyl)benzamide).

A food additive regulation is established permitting the combined residues of the insecticide flutolanil, N-(3-(1-methylethoxy)phenyl)-2-(trifluoromethyl)benzamide, and its metabolites converted to 2-(trifluoromethyl) benzoic acid and calculated as flutolanil in or on the following processed food commodity:

Commodity	Parts per million
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Commodity	Parts per million
Peanut meal .....	1.0

[FR Doc. 95-20015 Filed 8-15-95; 8:45 am]

BILLING CODE 6560-50-F

#### 40 CFR Parts 180 and 185

[OPP-300389A; FRL-4967-9]

RIN 2070-AB78

#### Sodium Propionate, Methoprene, and *Heliothis zea* NPV; Tolerance Actions

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Final rule.

**SUMMARY:** For each of the pesticides subject to the actions listed in this rule, EPA has completed the reregistration process and issued a Reregistration Eligibility Document (RED). In the reregistration process, all information to support a pesticide's continued registration is reviewed for adequacy and, when needed, supplemented with new scientific studies. Based on the RED tolerance assessments for the pesticide chemicals subject to this rule, EPA is taking the following tolerance actions: amending the exemptions from the requirement of a tolerance for methoprene; revoking exemptions for sodium propionate; and making wording changes to the exemption from the requirement of a tolerance for *Heliothis zea* NPV. With this rule to amend the exemptions from the requirement of tolerances for methoprene, the Agency is correcting its position in the RED, which stated that the exemptions should be revoked. The Agency believes that exemptions from the requirement of tolerances for these uses are appropriate.

**EFFECTIVE DATE:** This regulation becomes effective on August 16, 1995.

**ADDRESSES:** Written objections and hearing requests, identified by the document control number, [OPP-300389A], may be submitted to: Hearing Clerk (1900), Environmental Protection Agency, Rm. M3708, 401 M St., SW., Washington, DC 20460. Fees accompanying objections and hearing requests shall be labeled "Tolerance Petition Fees" and forwarded to: EPA Headquarters Accounting Operations Branch, OPP (Tolerance Fees), P.O. Box 360277M, Pittsburgh, PA 15251. A copy of any objections and hearing requests filed with the Hearing Clerk should be identified by the document control number and submitted to: Public

Response and Program Resources Branch, Field Operations Division (7506C), Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. In person, bring copy of objections and hearing requests to: Rm. 1132, CM #2, 1921 Jefferson Davis Hwy., Arlington, VA 22202.

A copy of objections and hearing requests filed with the Hearing Clerk may also be submitted electronically by sending electronic mail (e-mail) to: opp-docket@epamail.epa.gov. Copies of objections and hearing requests must be submitted as an ASCII file avoiding the use of special characters and any form of encryption. Copies of objections and hearing requests will also be accepted on disks in WordPerfect in 5.1 file format or ASCII file format. All copies of objections and hearing requests in electronic form must be identified by the docket number [OPP-300389A]. No Confidential Business Information (CBI) should be submitted through e-mail. Electronic copies of objections and hearing requests on this rule may be filed online at many Federal Depository Libraries. Additional information on electronic submissions can be found below in this document.

**FOR FURTHER INFORMATION CONTACT:** By mail: Philip Poli, Special Review and Reregistration Division (7505W), Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. Office location and telephone number: Crystal Station #1, 3rd Floor, 2800 Crystal Drive, Arlington, VA 22202, (703)-308-8038; e-mail: poli.philip@epamail.epa.gov.

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of June 28, 1995 (60 FR 33383), EPA issued a proposed rule (FRL-4960-5) affecting 40 CFR 180.2, 180.1015, 180.1027, 180.1033, and 185.4150 regarding various chemicals and tolerance actions the Agency proposed to take. Specifically, EPA proposed actions regarding the following chemicals: Methoprene, the revision of the methoprene regulation in 40 CFR 180.1033 to reflect changed uses and the revocation of the methoprene regulation in 40 CFR 185.4150; sodium propionate, the revocation of exemptions under 40 CFR 180.2(a) and 180.1015; and *Heliothis zea* NPV, the amendment of 40 CFR 180.1027 to better reflect the current viral identification and testing technology.

There were no comments or requests for referral to an advisory committee received in response to the proposed rule.

The data submitted with the proposal and other relevant material have been

evaluated and discussed in the proposed rule. Based on the data and information considered, the Agency concludes that the regulations issued in this document will protect the public health. Therefore, the regulations are established as set forth below.

Any person adversely affected by this regulation may, within 30 days after publication of this document in the **Federal Register**, file written objections and/or request a hearing with the Hearing Clerk, at the address given above (40 CFR 178.20). A copy of the objections and/or hearing requests filed with the Hearing Clerk should be submitted to the OPP docket for this rulemaking. The objections submitted must specify the provisions of the regulation deemed objectionable and the grounds for the objections (40 CFR 178.25). Each objection must be accompanied by the fee prescribed by 40 CFR 180.33(i). If a hearing is requested, the objections must include a statement of the factual issue(s) on which a hearing is requested, the requestor's contentions on such issues, and a summary of any evidence relied upon by the objector (40 CFR 178.27). A request for a hearing will be granted if the Administrator determines that the material submitted shows the following: There is a genuine and substantial issue of fact; there is a reasonable possibility that available evidence identified by the requestor would, if established, resolve one or more of such issues in favor of the requestor, taking into account uncontested claims or facts to the contrary; and resolution of the factual issue(s) in the manner sought by the requestor would be adequate to justify the action requested (40 CFR 178.32).

A record has been established for this rulemaking under docket number [OPP-300389A] (including any objections and hearing requests submitted electronically as described below). A public version of this record, including printed, paper versions of electronic comments, which does not include any information claimed as CBI, is available for inspection from 8 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The public record is located in Room 1132 of the Public Response and Program Resources Branch, Field Operations Division (7506C), Office of Pesticide Programs, Environmental Protection Agency, Crystal Mall #2, 1921 Jefferson Davis Highway, Arlington, VA.

Written objections and hearing requests, identified by the document control number [OPP-300389A], may be submitted to the Hearing Clerk (1900), Environmental Protection Agency, Rm.

3708, 401 M St., SW., Washington, DC 20460.

A copy of electronic objections and hearing requests filed with the Hearing Clerk can be sent directly to EPA at: opp-docket@epamail.epa.gov

A copy of electronic objections and hearing requests filed with the Hearing Clerk must be submitted as an ASCII file avoiding the use of special characters and any form of encryption.

The official record for this rulemaking, as well as the public version, as described above will be kept in paper form. Accordingly, EPA will transfer any objections and hearing requests received electronically into printed, paper form as they are received and will place the paper copies in the official rulemaking record which will also include all objections and hearing requests submitted directly in writing. The official rulemaking record is the paper record maintained at the address in "ADDRESSES" at the beginning of this document.

Under Executive Order 12866 (58 FR 51735, Oct. 4, 1993), the Agency must determine whether the regulatory action is "significant" and therefore subject to review by the Office of Management and Budget (OMB) and the requirements of the Executive Order. Under section 3(f), the order defines a "significant regulatory action" as an action that is likely to result in a rule (1) having an annual effect on the economy of \$100 million or more, or adversely and materially affecting a sector of the economy, productivity, competition, jobs, the environment, public health or safety, or State, local, or tribal governments or communities (also referred to as "economically significant"); (2) creating serious inconsistency or otherwise interfering with an action taken or planned by another agency; (3) materially altering the budgetary impacts of entitlement, grants, user fees, or loan programs or the rights and obligations of recipients thereof; or (4) raising novel legal or policy issues arising out of legal mandates, the President's priorities, or the principles set forth in this Executive Order.

Pursuant to the terms of the Executive Order, EPA has determined that this rule is not "significant" and is therefore not subject to OMB review.

Pursuant to the requirements of the Regulatory Flexibility Act (Pub. L. 96-354, 94 Stat. 1164, 5 U.S.C. 601-612), the Administrator has determined that regulations establishing new tolerances or raising tolerance levels or establishing exemptions from tolerance requirements do not have a significant

economic impact on a substantial number of small entities. A certification statement to this effect was published in the **Federal Register** of May 4, 1981 (46 FR 24950).

#### List of Subjects in 40 CFR Parts 180 and 185

Environmental protection, Administrative practice and procedure, Agricultural commodities, Food additives, Pesticides and pests, Reporting and recordkeeping requirements.

Dated: August 8, 1995.

**Lois Rossi,**

*Director, Special Review and Reregistration Division, Office of Pesticide Programs.*

Therefore, 40 CFR, chapter I, is amended as follows:

#### PART 180—[AMENDED]

1. The authority citation for part 180 continues to read as follows:

**Authority:** 21 U.S.C. 346a and 371.

2. Section 180.2 is revised to read as follows:

##### **§ 180.2 Pesticide chemicals considered safe.**

(a) As a general rule, pesticide chemicals other than benzaldehyde (when used as a bee repellent in the harvesting of honey), ferrous sulfate, lime, lime-sulfur, potassium carbonate, potassium polysulfide, potassium sorbate, sodium carbonate, sodium chloride, sodium hypochlorite, sodium polysulfide, sodium sesquicarbonate, sorbic acid, sulfur, and when used as plant desiccants, sodium metasilicate (not to exceed 4 percent by weight in aqueous solution) and when used as postharvest fungicide, citric acid, fumaric acid, oil of lemon, oil of orange, and sodium benzoate are not for the purposes of section 408(a) of the Act generally recognized as safe.

(b) Upon written request, the Registration Division will advise interested persons whether a pesticide chemical should be considered as poisonous or deleterious, or one not generally recognized by qualified experts, as safe.

(c) The training and experience necessary to qualify experts to evaluate the safety of pesticide chemicals for the purposes of section 408(a) of the Act are essentially the same as training and experience necessary to qualify experts to serve on advisory committees prescribed by section 408(g) of the Act. (See § 180.11.)

##### **§ 180.1015 [Removed]**

c. Section 180.1015 is removed.

d. Section 180.1027 is revised to read as follows:

##### **§ 180.1027 Nuclear polyhedrosis virus of *Heliothis zea*; exemption from the requirement of a tolerance.**

(a) For the purposes of this section, the viral insecticide must be produced with an unaltered and unadulterated inoculum of the single-embedded *Heliothis zea* nuclear polyhedrosis virus (HzSNPV). The identity of the seed virus must be assured by periodic checks.

(b) Each lot of active ingredient of the viral insecticide shall have the following specifications:

(1) The level of extraneous bacterial contamination of the final unformulated viral insecticide should not exceed  $10^7$  colonies per gram as determined by an aerobic plate on trypticase soy agar.

(2) Human pathogens, e.g., *Salmonella*, *Shigella*, or *Vibrio*, must be absent.

(3) Safety to mice as determined by an intraperitoneal injection study must be demonstrated.

(4) Identity of the viral product, as determined by the most sensitive and standardized analytical technique, e.g., restriction endonuclease and/or SDS-PAGE analysis, must be demonstrated.

(c) Exemptions from the requirement of a tolerance are established for the residues of the microbial insecticide *Heliothis zea* NPV, as specified in paragraphs (a) and (b) of this section, in or on all agricultural commodities including: corn, cottonseed, beans, lettuce, okra, peppers, sorghum, soybeans, and tomatoes.

e. Section 180.1033 is revised to read as follows:

##### **§ 180.1033 Methoprene; exemption from the requirement of a tolerance.**

Methoprene is exempt from the requirement of a tolerance in or on all raw agricultural commodities when used to control mosquito larvae including pastures, rice fields, vineyards, date palm orchards, nut orchards, berry orchards, and fruit orchards.

#### PART 185—[AMENDED]

2. In part 185:

a. The authority citation for part 185 continues to read as follows:

**Authority:** 21 U.S.C. 348.

b. Section 185.4150 is revised to read as follows:

##### **§ 185.4150 Methoprene.**

A tolerance of 10 parts per million is established for residues of isopropyl (E,E)-11-methoxy-3,7,11-trimethyl-2,4-

dodecadienoate) in or on the food additive commodity cereal grain milled fractions (except flour and rice hulls).

[FR Doc. 95–20305 Filed 8–15–95; 8:45 am]

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#### 40 CFR Parts 185 and 186

[PP 4H5683/R2156; FRL–4968–1]

RIN 2070–AB78

#### Hexazinone; Food/Feed Additive Regulations

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Final rule.

**SUMMARY:** This document establishes food and feed additive regulations for residues of the herbicide hexazinone (3-cyclohexyl-6-(dimethylamino)-1-methyl-1,3,5-triazine-2,4(1H,3H)-dione) and its metabolites (calculated as hexazinone) in sugarcane molasses. E.I. du Pont de Nemours & Co., Inc., petitioned for these regulations under the Federal Food, Drug and Cosmetic Act (FFDCA).

**EFFECTIVE DATE:** This regulation becomes effective August 16, 1995.

**ADDRESSES:** Written objections and hearing requests, identified by the document control number, [PP 4H5683/R2156], may be submitted to: Hearing Clerk (1900), Environmental Protection Agency, Rm. M3708, 401 M St., SW., Washington, DC 20460. Fees accompanying objections and hearing requests shall be labeled "Tolerance Petition Fees" and forwarded to: EPA Headquarters Accounting Operations Branch, OPP (Tolerance Fees), P.O. Box 360277M, Pittsburgh, PA 15251. A copy of any objections and hearing requests filed with the Hearing Clerk should be identified by the document control number and submitted to: Public Response and Program Resources Branch, Field Operations Division (7506C), Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. In person, bring copy of objections and hearing requests to: Rm. 1132, CM #2, 1921 Jefferson Davis Hwy., Arlington, VA 22202.

A copy of objections and hearing requests filed with the Hearing Clerk may also be submitted electronically by sending electronic mail (e-mail) to: opp-docket@epamail.epa.gov. Copies of objections and hearing requests must be submitted as an ASCII file avoiding the use of special characters and any form of encryption. Copies of objections and hearing requests will also be accepted